Department of Health & Human Services Centers for Medicare & Medicaid Services 61 Forsyth St., Suite 4T20 Atlanta, Georgia 30303-8909



November 18, 2009

Ms. Elizabeth A. Johnson Commissioner Cabinet for Health and Family Services Department of Medicaid Services 275 East Main Street, 6W-A Frankfort, Kentucky 40621-0001

Attention: Kevin Skeeters

RE: Kentucky Title XIX State Plan Amendment, Transmittal #08-014

Dear Ms. Johnson

We accept your request, dated November 16, 2009 to withdraw State Plan Amendment 08-014. We are returning the Form HCFA-179 and the proposed plan pages.

If you have any questions or need any further assistance, please contact Maria Donatto at (404) 562-3697 or Darlene Noonan at (404) 562-2707.

Sincerely,

Mary Kaye Justis, RN, MBA

Acting Associate Regional Administrator

Division of Medicaid & Children's Health Operations

Enclosures

DEPARTMENT OF HEALTH AND HUMAN SERVICES HEALTH CARE FINANCING ADMINISTRATION		
TRANSMITTAL AND NOTICE OF APPROVAL OF STATE PLAN MATERIAL	1. TRANSMITTAL NUMBER: 08-014	2. STATE Kentucky
FOR: HEALTH CARE FINANCING ADMINISTRATION	3. PROGRAM IDENTIFICATION: TITLE XIX OF THE SOCIAL SECURITY ACT (MEDICAID)	
TO: REGIONAL ADMINISTRATOR HEALTH CARE FINANCING ADMINISTRATION DEPARTMENT OF HEALTH AND HUMAN SERVICES	4. PROPOSED EFFECTIVE DATE January 1, 2009	
5. TYPE OF PLAN MATERIAL (Check One):		
☐ NEW STATE PLAN ☐ AMENDMENT TO BE COM	NSIDERED AS NEW PLAN	AMENDMENT
COMPLETE BLOCKS 6 THRU 10 IF THIS IS AN AME		amendment)
6. FEDERAL STATUTE/REGULATION CITATION: 42 USC Chapter 6A, Subchapter II, Part D, subpart vii, 256b; 42 USC 701(a)(2)	7. FEDERAL BUDGET IMPACT: a. FFY 2009 - savings of \$150,000 b. FFY 2010 - savings of \$200,000	
8. PAGE NUMBER OF THE PLAN SECTION OR ATTACHMENT:	9. PAGE NUMBER OF THE SUPERSEDED PLAN SECTION OR ATTACHMENT (If Applicable):	
Att. 4.19-B page 20.2		
	Same	
 10. SUBJECT OF AMENDMENT: This plan amendment reimburses 340B drug pricing entities a dispensing clotting factor. 11. GOVERNOR'S REVIEW (Check One): GOVERNOR'S OFFICE REPORTED NO COMMENT COMMENTS OF GOVERNOR'S OFFICE ENCLOSED NO REPLY RECEIVED WITHIN 45 DAYS OF SUBMITTAL 	X OTHER, AS SPECIFIED to Commissioner, Departs Services	D: Review delegated
12. SIGNATURE OF STATE AGENCY OFFICIAL:	16. RETURN TO:	
10 Tillet 1 . John	Department for Medicaid Services 275 East Main Street 6W-A Frankfort, Kentucky 40621	
13. TYPED NAME: Elizabeth A. Johnson		
14. TITLE: Commissioner, Department for Medicaid Services		
15. DATE SUBMITTED: October 21, 2008	-	
FOR REGIONAL OF		
17, DATE RECEIVED;	18. DATE APPROVED:	
PLAN APPROVED – ON 19. EFFECTIVE DATE OF APPROVED MATERIAL:	E COPY ATTACHED 20. SIGNATURE OF REGIONAL OF	PIQIAL: /
21. TYPED NAME:	I My chy	\mathcal{K}
	22. TITLE: fet hy mento	Kindhe
23. REMARKS:		i en en Albertan in en han en

Methods and Standards for Establishing Payment Rates — Other Types of Care

I. Drugs

A. Reimbursement

- 1. Participating pharmacies are reimbursed for the cost of the drug plus a dispensing fee. Payments shall not exceed the upper limits specified in 42 CFR 447.331 through 447.334.
- 2. Participating dispensing physicians are reimbursed for the cost of the drug only.
- 3. Providers will be reimbursed only for drugs supplied from pharmaceutical manufacturers who have signed a rebate agreement.
- 4. Comprehensive Hemophilia Diagnostic Treatment Centers (CHDTC) as defined by 42 USC Chapter 6A, Subchapter II, Part D, subpart vii, 256b(4)(g) are eligible for the 340B Drug Pricing Program and are the only entity in Kentucky that is currently participating in the program. The department shall reimburse a CHDTC a dispensing fee for hemophilia treatment, pharmaceutical replacement factors for clotting disorders, 340B drugs at twelve (12) and one-half (1/2) cents per unit. A unit is defined as enzymes that are dosed in terms of units of enzyme activity (as opposed to weight or volume) per kilogram of body weight.
- B. Payment Limits Payment for the cost of drugs shall be the lesser of:
 - 1. The Federal Maximum Allowable Cost (FMAC) of the drug for multiple source drugs other than those brand name drugs for which a prescriber has certified in writing as "brand medically necessary";
 - 2. The State Maximum Allowable Cost (SMAC). A SMAC may be established for any drug for which two or more A-rated therapeutically equivalent, multisource, non-innovator drugs with a significant cost difference exist. The SMAC will be determined taking into account drug price status (nonrebatable, rebatable), marketplace status (obsolete, regional availability), equivalency rating (A-rated), and relative comparable pricing. Other factors considered are clinical indications of generic substitution, utilization and availability in the marketplace. The source of comparable drug prices will be nationally recognized comprehensive data files maintained by a vendor under contract with the Department for Medicaid Services. Resources accessed to determine SMAC include Average Wholesale Price, Wholesale Acquisition Cost, and Direct Price (to retail pharmacies) with weights applied based on the distribution of the volume purchased.
 - a. Multiple drug pricing resources are utilized to determine the estimated acquisition cost for the generic drugs. These resources include pharmacy providers, wholesalers, drug file vendors such as First Data Bank, and pharmaceutical manufacturers;

TN No. <u>08-014</u> Supersedes TN No. <u>05-004</u>

Approval Date:

Effective Date: 1/1/2009

Methods and Standards for Establishing Payment Rates — Other Types of Care

C. Dispensing Fee

1. When establishing dispensing fees, the Department takes into consideration the conclusions of a report regarding the dispensing of prescription medications to persons eligible for Medicaid benefits. The report is based upon a survey of pharmacy dispensing costs in the Commonwealth of Kentucky, a review of academic literature, and the reimbursement rates of other payers. The report, required by state law, is submitted every three (3) years to the Governor and to the Legislative Research Commission. Utilizing the above information the Department establishes a reasonable dispensing fee.

Effective February 23, 2005, the dispensing fee for a generic drug prescription is \$5.00 and for a brand name drug prescription is \$4.50. The dispensing fee is applied to outpatient pharmacies and to long term care facilities.

- For nursing facility residents meeting Medicaid patient status, an incentive of two (2) cents
 per unit dose shall be paid to long term care pharmacists for repackaging a non-unit dose drug
 in unit dose form.
- 3. Comprehensive Hemophilia Diagnostic Treatment Centers (CHDTC) as defined by 42 USC Chapter 6A, Subchapter II, Part D, subpart vii, 256b(4)(g) are eligible for the 340B Drug Pricing Program and are the only entity in Kentucky that is currently participating in the program. The department shall reimburse a CHDTC a dispensing fee for hemophilia treatment, pharmaceutical replacement factors for clotting disorders, 340B drugs at twelve (12) and one-half (1/2) cents per unit. A unit is defined as enzymes that are dosed in terms of units of enzyme activity (as opposed to weight or volume) per kilogram of body weight.

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Approval Date: _____

Effective Date: 01/01/2009